

JS005580563A

United States Patent [19]

[54] MULTIPLE ANTIGEN PEPTIDE SYSTEM HAVING ADJUVANT PROPERTIES,

[76] Inventor: James P. Tam, 607 S. Wilson Blvd.,

331,489

Nashville, Tenn. 37215

May 3, 1993

Dec. 28, 1994

PCT/US93/04179

METHODS OF USE THEREOF

§ 102(e) Date: Dec. 28, 1994

PCT Pub. Date: Nov. 11, 1993

[87] PCT Pub. No.: WO93/22343

VACCINES PREPARED THEREFROM AND

Tam

[21] Appl. No.:

[22] PCT Filed:

[86] PCT No.:

§ 371 Date:

[11] Patent Number:

5,580,563

[45] **Date of Patent:**

Dec. 3, 1996

A. Smith et al, ESCOM, Leiden, pp. 845–846. Huang et al (1992) Ibid pp. 847–848.

Nardelli et al (1992) *Innovation and Perspectives in Solid Phase Synthesis*, Epton ed, Collected papers, Second International Symposium, 27th–31st. Aug., 1991, Canterbury, England, Intercept–Limited, Andrew, pp. 241–249.

Chem. Abstract. 118(13) 29 Mar. 1993 #125070j on p. 869 of Nardelli et al. (1992) *Innovation and Perspectives in Solid Phase Synthesis*, Epton ed. Collected pprs. 2nd Int. Symp. Aug. 27–31/1991, Canterbury, Engl; Intercept Limited, Andover pp. 241–249.

Defoort et al (1992) Proc. Natl. Acad. Sci. USA 89(9): 3879-83.

Defoort et al (1992) International J. Peptide and Protein Res. 80 (3/4) pp. 214–221.

Primary Examiner—Kay K. A. Kim Attorney, Agent, or Firm—Klauber & Jackson

Related U.S. Application Data

[63] Continuation-in-part of Ser. No. 877,613, May 1, 1992, abandoned.

[51] **Int. Cl.**⁶ **A61K 39/39**; A61K 39/385; A61K 9/127; A61K 39/02

[52] **U.S. Cl.** **424/197.11**; 424/193.1; 424/278.1; 424/DIG. 16; 424/450; 424/196.11; 424/194.1; 530/345; 530/403

[56] References Cited

PUBLICATIONS

Defoort et al (1992) in *Peptides Chemistry and Biology* Proceedings of the Twelfth American Peptide Symposium Jun. 16–21, 1991, Cambridge, Massachussets, USA, ed by J.

[57] ABSTRACT

A multiple antigenic peptide system is disclosed that comprises a dendritic core and peptide and a lipophilic anchoring moiety. This particular combination has as an advantage that it eliminates the need for the inclusion of adjuvants found to be toxic to humans, and facilitates the exponential amplification of the antigenic potential of a vaccine prepared therefrom, as noncovalent amplification by a liposome or micellar form is possible. Further, multiple different antigenic peptides may be attached so that the system may be prepared for administration to concurrently treat diverse ailments, such as for example, AIDS and influenza. The present multiple antigen peptide system is capable of eliciting an immune response when injected into a mammal, and accordingly, vaccines prepared from the system and methods of use including therapeutic protocols are included.

36 Claims, 12 Drawing Sheets

